



You don't have to wait until your baby is born to learn more about their health.

Get crucial insights as early as nine weeks into your pregnancy with MaterniT[®] GENOME, a noninvasive prenatal screening (NIPS/NIPT) test.




Call Us


Toll-free (within the US): 877-821-7266
Billing/Cost Questions: 844-799-3243

Our Every Mom Pledge team is ready to answer questions about your insurance coverage and cost options.

womenshealth.labcorp.com/estimatemycost

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References

1. Rafalko, J., Soster, E., Caldwell, S. et al. Genome-wide cell-free DNA screening: a focus on copy-number variants. *Genet Med* 23, 1847–1853 (2021). <https://doi.org/10.1038/s41436-021-01227-5>



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PATIENT INFORMATION

MaterniT[®] Genome NIPS (NIPT)

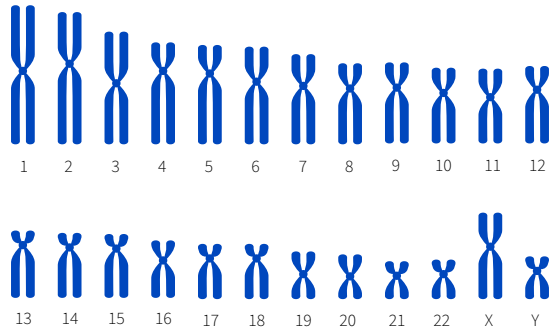
Every chromosome plays a role in your baby's development



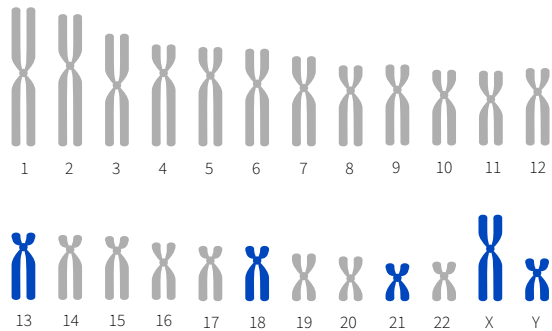
Why does every chromosome matter?

Most noninvasive prenatal screening (NIPS/NIPT) tests provide information on select chromosomes, but differences can be found in all chromosomes—which is why MaterniT GENOME covers them all.

Whole chromosomes analyzed by MaterniT GENOME



Whole chromosomes analyzed by most NIPSS (NIPTs)



Visit our website for additional information and online genetics resources.

What makes MaterniT GENOME different?

We found that up to 25% of all NIPS (NIPT) results could be detected only by MaterniT GENOME.¹ Most NIPSS (NIPTs) don't look for that 25%, but not looking doesn't mean they're not there.¹

What will MaterniT GENOME tell me?

Like most NIPSS (NIPTs), MaterniT GENOME can tell you if you have a higher or lower risk for trisomies 21 (Down syndrome), 18 (Edwards syndrome), and 13 (Patau syndrome). It can also predict if you're more likely to have a boy or a girl, and can identify other chromosomal changes, some of which may go undiagnosed at birth. Early detection can mean better care for you and your baby, before and after delivery.

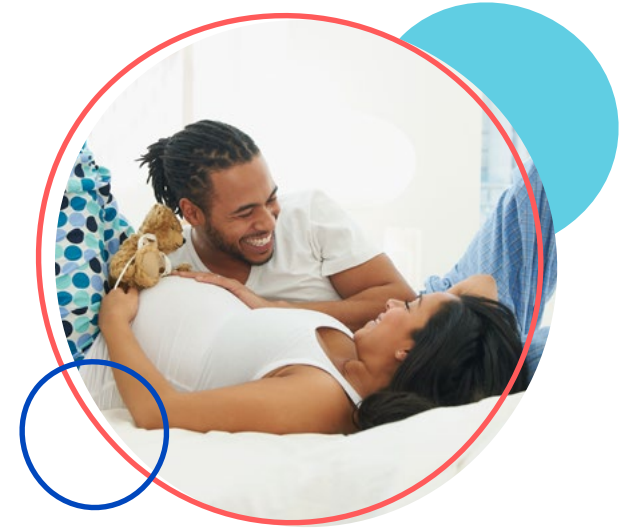
MaterniT GENOME is a screening test, and will deliver a result indicating whether there is increased or decreased risk for the conditions being screened. And like many screening tests, there is a risk of false negative and false positive results. Only a diagnostic test will deliver a definitive positive or negative result, so remember to speak with your healthcare provider about your test results.

MaterniT GENOME reports on:

Any trisomy or monosomy	<ul style="list-style-type: none"> Trisomy: an extra copy of a chromosome is present (3 instead of 2) Monosomy: a missing copy of a chromosome (1 instead of 2)
Sex chromosome abnormalities	<ul style="list-style-type: none"> An atypical number of X and/or Y chromosomes beyond typical female (XX) or male (XY) complement
Partial chromosome abnormalities	<ul style="list-style-type: none"> Missing or extra parts of the chromosome

Faster, clearer results for every parent

You have questions, and you want answers that are easy to understand. With MaterniT GENOME, you'll receive results, typically within five days of our lab receiving your sample.



Pioneering science, personalized service

Understand your cost options

Visit our website for your cost estimate and personalized payment options. Learn about our Moms Helping Moms of Tomorrow initiative.

Convenient blood draws

Getting your blood drawn is easier than ever. We have a nationwide network of patient service centers, allowing for convenient access to sample collection. Visit [Labcorp.com](https://www.labcorp.com) to find your nearest location.

Genetic counseling and support

Results may lead to more questions. Labcorp offers virtual results genetic counseling from a nationwide network of genetic counselors. You can also explore additional genetics resources on our website: [womenshealth.labcorp.com/patients/online-genetics-resources](https://www.womenshealth.labcorp.com/patients/online-genetics-resources)

Every Mom Pledge

We believe every mom should have access to the best possible care. That's why we work directly with you to make sure our testing services are accessible and out-of-pocket costs are transparent.